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By Fax

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Representative Michael Bilirakis
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Representative Sherrod Brown
Ranking Member
House Committee on Energy and
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2438 Rayburn House Office Building
Washington, D.C. 20515

Re: Pharmaceutical Pricing and Legislative Reform

Dear Senators Gregg and Kennedy, and Representatives Bilirakis and Brown:

Access to affordable prescription drugs is a critical issue that commands bipartisan support among State Attorneys General. The high price of brand name prescription drugs makes many life-saving medications unobtainable for large sectors of our population. This most adversely affects the less fortunate residents of our states, especially the uninsured, underinsured and senior citizens.

The cost of a brand name drug is greatly reduced when its generic equivalent is introduced into the market. We expect that in the coming years, generic alternatives to brand name pharmaceuticals will substantially increase consumers' access to necessary pharmaceutical care. Generic alternatives will also reduce the States' Medicaid costs for prescription drug reimbursements.

Since 1998, the Attorneys General have prosecuted numerous antitrust violations against brand name and generic pharmaceutical companies.¹ These companies have used their pharmaceutical patents to deter and foreclose generic competition. During the course of our investigations, we have observed a number of anticompetitive actions that continue to alarm us. These include:

- Improper Use of the Courts by brand name manufacturers, designed to delay entry by generic competitors under the 30-month stay provision of the Hatch-Waxman Act and to invoke antitrust immunity under federal antitrust laws.
- Patent Stacking and Fraud upon the Patent & Trademark Office ("PTO") and the Food & Drug Administration ("FDA"), in order to acquire multiple patents and improper patents, and to improperly list patents for the sole purpose of triggering 30-month stays on FDA approval of generic equivalents.
- **Anticompetitive agreements** between brand name manufacturers and their generic competitors to delay or foreclose entry of generic equivalents.
- Collusion between generic companies, who manipulate their 180-day marketing exclusivity under the Hatch-Waxman Act instead of bringing their products to market expeditiously.
- False and Misleading Price Reporting by certain generic manufacturers resulting in Medicaid over-reimbursements to pharmacists and physicians.

We support the safe and efficient availability of generic alternatives to brand name drugs, while protecting the valid intellectual property rights of the brand name drug companies. Therefore we, the undersigned Attorneys General, urge Congress to carefully examine the relationship between rising prescription drug costs and the above referenced anticompetitive behavior, and to enact legislation preserving and promoting generic competition for prescription drugs.

Attorney General G. Steven Rowe Attorney General of Maine

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¹ Some of these cases and pending settlements include: In re Cardizem CD Antitrust Litigation, MDL Docket No. 1278 (E.D. Mich), In re Terazosin Hydrochloride Antitrust Litigation, MDL Docket No. 1317 (S.D. Fla.), In re Buspirone Antitrust Litigation, MDL Docket No. 1413 (S.D.N.Y.) and State of Ohio, et al v. Bristol-Myers Squibb, Co., 1:02-cv-01080 (EGS) (D.D.C.).

² PPTF is the abbreviation for NAAG's Pharmaceutical Pricing Task Force.

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cc: Senator Bill Frist - Majority Leader Senator Tom Daschle - Minority Leader Representative J. Dennis Hastert - Speaker of the House Representative Nancy Pelosi - Minority Leader

Members of the Senate Committee on the Judiciary's Subcommittee on Antitrust, Competition and Business and Consumer Rights

Members of the Senate Committee on Health, Education, Labor and Pensions

Members of the House Committee on Energy and Commerce's Subcommittee on Health

Members of the House Committee on the Judiciary's Subcommittee on Courts, the Internet, and Intellectual Property